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(71) Applicant: **Bristol-Myers Squibb Company
345 Park Avenue
New York, N.Y. 10154(US)**

(72) Inventor: **Soichiro, Toda
3-591-5, Ohnari-cho
Ohmiya(JP)**

Inventor: **Toshikazu, Oki
4-20-1 Shodo, Sakae-ku
Yokohama(JP)**

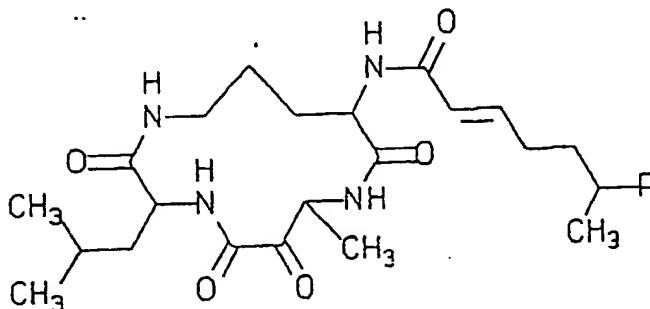
Inventor: **Yasutaro, Hamagishi
2-12-34, Tsujido-shinmachi
Fujisawa(JP)**

Inventor: **Koji, Tomita
5-2-3, Kami-Yoga
Setagaya-ku, Tokyo(JP)**

(74) Representative: **Kinzebach, Werner, Dr. et al
Patentanwälte Reitschötter, Kinzebach und
Partner Sternwartstrasse 4 Postfach 86 06 49
W-8000 München 86(DE)**

(54) **1,4,8-Triazacyclotridecane derivatives having prolyl endopeptidase inhibiting activity.**

(57) Disclosed herein are antibiotic BU-4164E A and B, having formule (1), isolated from Streptomyces sp. These compounds are prolyl endopeptidase inhibitors.



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The present invention relates to novel compounds designated BU-4164E A and B, which are produced from Strain No. R353-21, isolated from a soil sample collected in India. Based on the morphology, cell chemistry and cultural and physiological characteristics, Strain No. R353-21 was identified as Streptomyces sp.

- 5 BU-4164E A and B are prolyl endopeptidase (PED) inhibitors which have anti-amnestic and/or anti-dementia properties. Physico-chemical properties and structural studies indicate that these compounds are novel 13 membered cyclic peptides containing UV-absorbing chromophore.

- Prolyl endopeptidase, a serine protease, plays an important role in the degradation of biologically active peptides such as vasopressin, oxytocin, angiotensins I and II, substance P, bradykinin, neurotensin, etc. 10 Prolyl endopeptidase hydrolyzes these peptides and regulates their biological activities. D. De Wied and coworkers previously suggested and recently gave some evidence that vasopressin was involved in the process of learning and memory of animals.¹⁻⁴ In addition, the Nippon Roche group found an anti-amnestic agent aniracetam to inhibit PED.⁵ T. Yoshimoto (Nagasaki Univ.) synthesized specific PED inhibitors such as Z-Gly-Pro-CH₂Cl, Z-Pro-prolinal and Z-Val-prolinal based on the specificity of substrate for this enzyme and demonstrated correlation between their anti-amnestic activity in rats and PED inhibitory activity.⁶ H. Kaneto, et al. also confirmed anti-amnestic activity of vasopressin and the above synthetic PED inhibitors using several mouse models.⁷

U.S. Patent Nos. 4,857,537, 4,857,524, 4,826,870, 4,810,721, 4,772,587, 4,757,083, 4,743,616, and 4,701,465 relate to the synthetic production of compounds which are PED inhibitors.

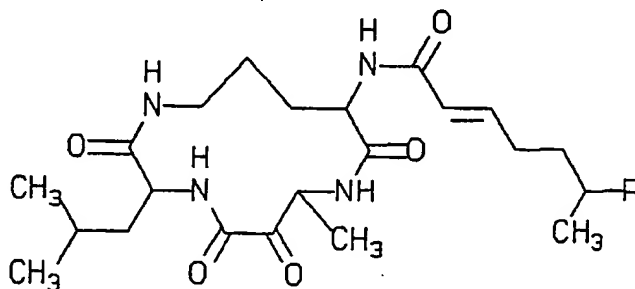
- 20 Taking into consideration the above information, PED inhibitor screening was initiated by the present inventors with an intention to discover new anti-amnestic and/or anti-dementia compounds from microbial products.

The present invention provides the novel PED inhibitors and anti-amnestic and/or anti-dementia compounds designated BU-4164E A and B which compounds have the following formula:

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wherein R is from C₁-C₄ alkyl, preferably methyl or ethyl.

- Another aspect of the invention provides a process for the production of BU-4164E A and B, which 40 comprises culturing an strain of Streptomyces sp. preferably Strain No. R353-21.

In another aspect of the invention there are provided pharmaceutical compositions useful for treating amnesia and/or dementia in a mammalian host comprising an effective amount of BU-4164E A and B together with a pharmaceutically acceptable carrier.

Figure 1 shows the IR spectra of BU-4164E A.

- 45 Figure 2 shows the IR spectra of BU-4164E B.

Figure 3 shows the ¹H-NMR spectra of BU-4164E A.

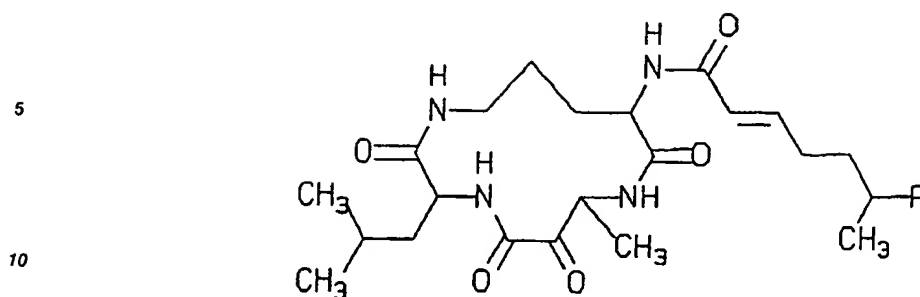
Figure 4 shows the ¹H-NMR spectra of BU-4164E B.

Figure 5 shows the ¹³C-NMR spectra of BU-4164E A.

Figure 6 shows the ¹³C-NMR spectra of BU-4164E B.

- 50 The BU-4164E A and B of the present invention have the following formula:

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wherein R is from C₁-C₄ alkyl, preferably methyl or ethyl.

15 These compounds are produced by fermentation of a BU-4164E A and B-producing strain of *Streptomyces* sp. The active principle was recovered from the cultured broth by solvent extraction and purified by column chromatography to obtain two components, BU-4164E A and B. These compounds demonstrated a high level of inhibition activity against a rabbit brain PED and *Flavobacterium meningosepticum* PED. *Flavobacterium* PED was inhibited with IC₅₀: 0.0021 - 0.0037 µg/ml, other proteases tested were not
20 inhibited.

The preferred producing organism is a novel strain of a known species of *Streptomyces* designated herein as *Streptomyces* sp., strain no. R353-21. This strain was isolated from a soil sample collected in India. A biologically pure culture of this strain was deposited with the American Type Culture Collection, Rockville, Maryland, under the accession number ATCC 55001.

25 **Morphology** Both substrate and aerial mycelia are formed. Spore-chains are born monopodially or often in tuft on the aerial mycelium, and are open irregular spiral, loop or straight. The chains contain 10 to 30 or more spores per chain. The spore chain morphology belongs to *Retinaculum-Apertum*. Scanning electron microscopy indicates that the spores are spherical to oval (0.7-1.0 x 0.7-1.5 µm), and have a smooth surface.

30 **Cultural and physiological characteristics** The color of aerial mycelium is gray with shades of yellow, brown, olive or green. The substrate mycelium colors to dark reddish brown in ISP medium Nos. 2, 5 and 7, and dark olive brown in ISP medium No. 3. These pigments are more or less diffusible. Melanin is produced in ISP No. 1 broth and ISP No. 6 agar.

Gelatin and starch are hydrolyzed. Skim-milk is coagulated and peptonized. Nitrate is reduced to nitrite in Czapek's nitrate broth but not in peptone broth. Tyrosinase is produced. Growth occurs in 7% NaCl agar but not in 9%. Growth temperature ranges between 13°C and 50°C. No growth is seen at 10°C and 52°C.

The cultural characteristics and the utilization of sugars are shown in Tables 1 and 2, respectively.

Cell chemistry Whole cell hydrolysate contains LL-diaminopimelic acid.

40 **Taxonomic position** The above-mentioned morphology, cultural and physiological characteristics and cell chemistry of strain R353-21 indicate that the strain belongs to the genus *Streptomyces*.

According to the descriptions of Pridham and Tresner^a and Shirling and Gottlieb,⁹ strain R353-21 resembles many known species of the Genus, particularly *Streptomyces ambofaciens*, *S. eurythermus*, *S. griseosporus*, *S. luteogriseus*, *S. microflavus*, *S. neyagawaensis*, *S. olivochromogenes*, *S. resistomycificus*, and *S. violaceochromogenes*. To determine the species of strain R353-21, additional comparative studies
45 are necessary. At present, strain R353-21 was designated as a new strain of *Streptomyces* sp.

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Table 1. Cultural characteristics of strain R353-21

Medium	Growth	Aerial mycelium	Substrate mycelium	Diffusible pigment
Sucrose-nitrate agar (Czapek-Dox agar)	Scant	No or trace	Colorless	Pale yellow (89)
Tryptone-yeast extract broth (ISP No. 1)	Good; pellicle not turbid	No	Colorless	Deep brown (56)
Yeast extract-malt extract agar (ISP No. 2)	Good	Abundant; greenish gray (155)	Dark reddish brown (44)	Grayish reddish brown (46)
Oatmeal agar (ISP No. 3)	Good	Abundant; olive gray (113)	Dark olive brown (96)	Dark grayish yellow (91)
Inorganic salts-starch agar (ISP No. 4)	Moderate	Moderate; medium gray (265)	Dark yellowish brown (78)	Grayish yellow (90)
Glycerol-asparagine agar (ISP No. 5)	Good	Abundant; light brownish gray (63)	Moderate reddish brown (43)	Moderate reddish brown (43)
Peptone-yeast extract- iron agar (ISP No. 6)	Moderate	No or trace	Black	Black
Tyrosine agar (ISP No. 7)	Good	Abundant; light brownish gray (63)	Dark grayish reddish brown (47)	Grayish brown (61)
Glucose-asparagine agar	Moderate	Moderate; light brownish gray (63)	Strong brown (55)	Brownish orange (54)
Bennett's agar	Good	Abundant; dark grayish yellow (91)	Dark reddish brown (44)	Grayish reddish brown (46)
Nutrient agar	Poor	No or trace	Moderate yellowish brown (77)	Light yellowish brown (76)

Observation after incubation at 28°C for 2 weeks
Color name, used: ISCC-NBS color-name charts

Table 2. Sugar utilization of strain R353-21

5	Glycerol	+	Cellobiose	+
	D-Arabinose	-	Melibiose	+
	L-Arabinose	+	Trehalose	+
10	D-Xylose	+	Raffinose	+(w)
	D-Ribose	+	D-Melezitose	-
15	L-Rhamnose	+(w)	Soluble starch	+
	D-Glucose	+	Cellulose	-
	D-Galactose	+	Dulcitol	-
20	D-Fructose	+	Inositol	+(w)
	D-Mannose	+	D-Mannitol	+
25	L-Sorbose	-	D-Sorbitol	-
	Sucrose	+(w)	Salicin	+
30	Lactose	+		

Basal medium: Pridham-Gottlieb's inorganic medium
(= ISP medium No. 9)

+(w), weakly positive

BU-4164E A and B may be produced by cultivating a BU-4164E A and B-producing strain of *Streptomyces* sp., ATCC 55001 or a mutant or variant thereof, under submerged aerobic conditions in an aqueous nutrient medium. The organism is grown in a nutrient medium containing an assimilable carbon source, for example an assimilable carbohydrate. Examples of suitable carbon sources include lactose, glycerol, sucrose, corn starch, glucose, mannose, fructose, cellobiose, trehalose, mannitol and xylose. The nutrient medium should also contain an assimilable nitrogen source such as, for example, fish meal, peptone, soybean flour, peanut meal, cotton seed meal and corn steep liquor. Nutrient inorganic salts may also be incorporated in the medium and such salts may comprise any of the usual salts capable of providing sodium, potassium, ammonium, calcium, phosphate, sulfate, chloride, bromide, nitrate, carbonate or like ions.

Production of the BU-4164E A and B compounds may be effected at any temperature conducive to satisfactory growth of the organism, i.e., approximately 14-44°C and is conveniently carried out at a temperature of about 28°C. Ordinarily, optimum production is obtained after incubation periods of about 3-6 days. The fermentation may be carried out in flasks and in laboratory or industrial fermentors of various capacities. When tank fermentation is to be carried out, it is desirable to produce a vegetative inoculum in a nutrient broth by inoculating the broth culture with a slant or soil culture or a lyophilized culture of the

organism. The medium in which the vegetative inoculum is produced can be the same as, or different from, that utilized in the tank for the production of the new compounds of the present invention as long as it is such that a good growth of the microorganism is obtained.

Production of BU-4164E A and B can be followed during the fermentation by testing samples of the
 5 broth or extracts of the mycelial solids for activity against organisms known to be sensitive to the compounds of the present invention or by an in vitro cytotoxicity assay, e.g. using rabbit brain PED and/or Flavobacterium meningosepticum PED.

When fermentation is complete, BU-4164E A and B are recovered from the fermentation broth and separated by extraction with a suitable organic solvent followed by a series of column chromatographies.
 10 Examples 1 and 2 below illustrate specific procedures for obtaining BU-4164E A and B in substantially purified form.

As is the case with other microorganisms, the characteristics of the new BU-4164E A and B-producing strain of the present invention, Streptomyces sp., ATCC 55001, are subject to variation. Recombinants, variants and mutants of the ATCC 55001 Strain, may be obtained by treatment with various known
 15 mutagens such as ultraviolet rays, X-rays, high frequency waves, radioactive rays and chemicals. Natural and induced variants, mutants and recombinants of Streptomyces sp., ATCC 55001 which retain the characteristic of producing BU-4164E A and B are intended to be encompassed by the present invention.

Physico-chemical properties of BU-4164E A and B

20 BU-4164E A and B were obtained as white amorphous powder. Their physico-chemical properties are very similar to each other as summarized in Table 3. They are soluble in dimethyl sulfoxide but practically insoluble in other organic solvents and water. BU-4164E A and B gave negative response to ninhydrin, Tollens and Sakaguchi reactions. The IR spectra of BU-4164E A and B (Fig. 1 and 2, respectively) possess
 25 strong absorption at 1660 and 1520 - 1540 cm^{-1} , suggesting that they come under the peptide group of antibiotics. In the UV spectra, BU-4164E A and B showed absorption maxima at 212 and 213 nm, respectively. The ^1H -NMR spectra and the ^{13}C -NMR spectra of BU-4164E A and B are shown in Fig. 3 and 4 and Fig. 5 and 6, respectively. Complete acid hydrolysis of both components gave the same ninhydrin-
 30 positive products, two of which were identified as ornithine and leucine by the amino acid analysis.

Table 3. Physico-chemical properties of BU-4164E A and B.

		BU-4164E A	BU-4164E B
5	Nature	White Powder	White Powder
	M.P.	>280°C (dec.)	>280°C (dec.)
10	$[\alpha]_D^{26}$ (C=0.25, DMSO)	-134°	-96°
	MS(SI-MS)M/Z)	451 (M+H) ⁺	465 (M+H) ⁺
	Molecular Formula	C ₂₃ H ₃₈ N ₄ O ₅	C ₂₄ H ₄₀ N ₄ O ₅
15	UVλ _{max} ^{50%MeOH_{nm}} (E _{1%} ^{1cm})	212 (767)	213 (645)
20	Color Reaction	Ninhydrin(-), Sakaguchi(-), Tollens(-)	
	Solubility	Soluble in DMSO	Soluble in DMSO
	Degradation Products	Ornithine	Ornithine
25		Leucine	Leucine
		$\begin{array}{c} \text{CH}_3-\text{CH}-\text{C}-\text{COOH}^* \\ \quad \parallel \\ \text{NH}_2 \quad \text{O} \end{array}$	$\begin{array}{c} \text{CH}_3-\text{CH}-\text{C}-\text{COOH}^* \\ \quad \parallel \\ \text{NH}_2 \quad \text{O} \end{array}$
30		C ₅ H ₁₁ CH=CHCOOH	C ₆ H ₁₃ CH=CHCOOH

*Isolated as $\begin{array}{c} \text{CH}_3\text{CH}-\text{CHCOOH} \\ | \\ \text{NH}_2\text{OH} \end{array}$ after NaBH₄ reduction followed by acid

hydrolysis

According to one aspect of the invention, therefore, there is provided a method for therapeutically treating a mammalian host affected by amnesia and/or dementia which comprises administering to said host an effective amount of BU-4164E A or B, or combinations thereof.

In yet another aspect of this invention a pharmaceutical composition is provided which comprises an effective amount of BU-4164E A or B or combinations thereof, in combination with an inert pharmaceutical acceptable carrier or diluent. These compositions can be made up in any pharmaceutical form appropriate for the desired route of administration.

The pharmaceutical compositions provided by the present invention may contain other active ingredients, e.g. other anti-amnesic and/or anti-dementia agents, and may be made up in any form appropriate for the desired route of administration. Examples of such compositions include solid compositions for oral administration such as capsules, tablets, pills, powders and granules, liquid compositions for oral administration such as solutions, suspensions, syrups or elixirs and preparations for parenteral administration such as sterile solutions, suspensions or emulsions. They may also be manufactured in the form of sterile solid compositions which can be dissolved in sterile water, physiological saline or other sterile injectable medium immediately before use.

The dosage ranges will commonly range from about 50-200 mg, but actual amounts will vary according to the particular compound being used, the particular composition formulated, the mode of administration and the particular situs, host and disease being treated. Many factors that modify the action of the drug will be taken into account by those skilled in the art, e.g. age, body weight, sex, diet, time of administration, route of administration, rate of excretion, condition of the host, drug combinations, reaction sensitivities and severity of the disease. Optimal dosages for a given set of conditions can be ascertained by those skilled in

the art using conventional dosage determination tests in view of the experimental animal data provided.

While the invention has been described with respect to various specified examples and embodiments, it is to be understood that the invention is not limited thereto.

5 EXAMPLE 1

Fermentation

A loopful of the slant culture of *Streptomyces* sp. strain No. R353-21 which matured on modified
 10 Bennett's medium was inoculated into a 500-ml Erlenmeyer flask containing 100 ml of the vegetative medium consisted of soluble starch (Nichiden Kagaku) 3%, Bacto-liver (Difco) 1%, polypeptone (Daigo Eiyo Kagaku) 0.5%, NaCl 0.3%, $(\text{NH}_4)_2\text{SO}_4$ 0.1% and CaCO_3 0.6%, pH being adjusted to 7.0 before autoclaving. The culture was incubated at or about 28 °C for 3 days on a rotary shaker (200 rpm) and 5 ml of the growth
 15 was transferred to a 500-ml Erlenmeyer flask containing 100 ml of the production medium having the same composition as the above vegetative medium. The fermentation was carried out at or about 28 °C for 4 days on a rotary shaker (200 rpm). The antibiotic production in the fermentation broth was monitored by the prolyl endopeptidase inhibition assay method. The fermentation broth showed a maximum inhibition activity in terms of IC_{50} at x640 dilution unit after 3 day-incubation.

For a large scale production, a tank fermentation was carried out. Two liters of the seed culture
 20 prepared by flask fermentation was transferred to a 200-L tank fermentor containing 120 liters of the production medium having the same composition as described before. The fermentation was carried out at or about 28 °C for 3 days with agitation at 250 rpm and air flow rate at 120 liters per minute.

EXAMPLE 2

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Extraction and purification

The harvested broth (60 L), obtained by the general procedure of Example 1, was vigorously stirred
 with n-butanol (32 L) for about 1 hour. The organic layer (25 L) was separated by the aid of a Sharples
 30 centrifuge (Kokusan No. 4A) and evaporated to dryness in vacuo. The residue was shaken with a mixture of water (2.3 L) and ethyl acetate (2.5 L). The ethyl acetate layer was washed with alkaline water (pH 9.0, 2.5 L) and acidic water (pH 2.0, 2.5 L), and then concentrated to dryness in vacuo to give brown oil (20.1 g, IC_{50} vs *Flavobacterium* PED: 1.25 $\mu\text{g/ml}$). The oil was applied to a column of Diaion HP-20 (Mitsubishi Chem. Ind. Tokyo, 800 ml). The column was developed with water (2 L), 50% methanol (2 L), 60%
 35 methanol (2 L), 70% methanol (1.5 L) and 80% methanol (2 L), successively. The eluate after 70% methanol elution was collected in 15 ml-fractions and each fraction was determined its PED activity. The combined active fractions (No. 97 - 170) were concentrated in vacuo to give 2.15 g of brown powder (IC_{50} : 0.13 $\mu\text{g/ml}$). This sample was chromatographed on a reversed phase silica gel column (YMC-ODS, 800 ml) eluting with 30% (15 ml-cut, fraction No. 1 - 50), 35% (No. 51 - 100), 40% (No. 101 - 150) and 45% (No.
 40 151-) acetonitrile in 0.15M phosphate buffer, pH 3.5. The PED active fractions (No. 63 - 155) were collected and concentrated to an aqueous solution. The concentrate was then extracted with n-butanol and the extract was evaporated in vacuo to give a crude solid containing BU-4164E A and B (171 mg, IC_{50} : 0.015 $\mu\text{g/ml}$). The mixture (40 mg) was purified by a reversed phase silica gel column chromatography (YMC-ODS, 150 ml). The column was washed with 30% acetonitrile in 0.15M phosphate buffer (pH 7.0, 500 ml) and then
 45 developed with 35% acetonitrile solution (fraction 15 ml-cut). Fractions No. 10 - 15 were combined and concentrated to an aqueous solution, which was extracted with n-butanol. The extract was evaporated in vacuo to dryness. The residue was chromatographed on Sephadex LH-20 (ϕ 1 x 35 cm) with dichloromethane-methanol (1:2) elution to give 14 mg of BU-4164E A. Fractions No. 26 - 33 in the above reversed phase silica gel chromatography were similarly treated to give 9 mg of BU-4164E B.

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EXAMPLE 3

Biological activity of BU-4164E A and B: Prolyl endopeptidase inhibition

55 Two enzymes were used: *Flavobacterial* enzyme for the screening and the isolation study and partially purified rabbit brain enzyme for *in vitro* evaluation. Ten μl of *Flavobacterium meningosepticum* PED, ¹⁰, ¹¹ (Seikagaku Kogyo Co. Ltd., 0.5 unit/ml in 50mM phosphate buffer, pH 7.0) or rabbit brain PED partially purified from the whole brain homogenate¹², 13.5 μl of a test sample and 86.5 μl of 100mM phosphate

buffer (pH 7.0) were mixed in a well of a 96-well microplate and pre-incubated for about 30 minutes at about 30° C. The reaction was started by addition of 25 µl of 2mM carbobenzyloxyglycyl-prolyl-p-nitroanilide (Z-Gly-Pro-pNA) in 40% dioxane-100mM phosphate buffer, pH 7.0. After about 5 minute-incubation at about 30° C, the amount of released p-nitroaniline was determined colorimetrically with a microplate autoreader (Titertek Multiskan^R MMC) at 414 nm. For control, the reaction buffer was added to the reaction mixture instead of the test sample. The percentage of inhibition was calculated by the following equation:

$$\% \text{ Inhibition} = \frac{\text{Abs}_{414\text{nm}} (\text{Control}) - \text{Abs}_{414\text{nm}} (\text{Sample})}{\text{Abs}_{414\text{nm}} (\text{Control})} \times 100$$

Activity of BU-4164E A and B and the reference compound, Z-Val-prolinal is summarized in Table 5.

Table 5. PED Inhibitory activities of BU-4164E A and B

<u>IC₅₀ (µg/ml)</u>			
<u>PED</u>			
<u>Compound</u>	<u>Flavobacterium</u>	<u>Rabbit brain</u>	
BU-4164E A	0.0037	0.085	
BU-4164E B	0.0021	0.031	
Z-Val-prolinal	0.0036	0.016	

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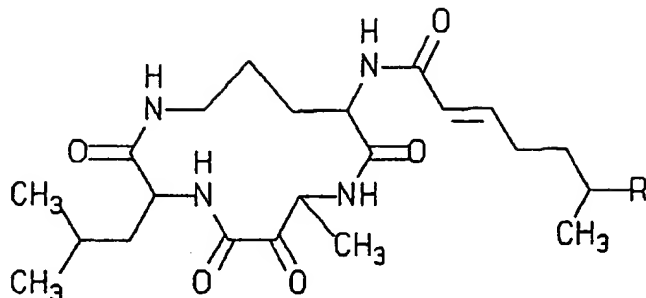
Claims

1. A compound having the structure

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wherein R is from C₁-C₄ alkyl.

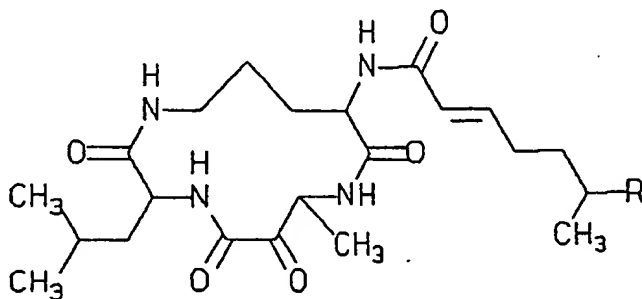
2. The compound of Claim 1 wherein R is methyl or ethyl.
3. A process for producing a compound of claims 1 or 2 which comprises cultivating a BU-4164E A or B-producing strain of *Streptomyces* sp. or mutant or variant thereof, under submerged aerobic conditions in an aqueous nutrient medium containing assimilable sources of carbon and nitrogen until a substantial amount of BU-4164E A or B is produced by said organism in said culture medium and then recovering the BU-4164E A or B from the culture medium.
4. A biologically pure culture of the microorganism *Streptomyces* sp., ATCC 55001, or a variant or mutant thereof, capable of producing the antibiotic BU-4164E A or B upon cultivation in an aqueous nutrient medium.
5. A pharmaceutical composition comprising an effective amount of at least one compound of claims 1 or 2, in combination with an inert pharmaceutically acceptable carrier or diluent.
6. The use of at least one compound according to claims 1 or 2 for preparing a pharmaceutical composition for therapeutically treating amnesia or dementia.

Claims for the following Contracting State: ES

1. A process for producing a compound having the structure

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- wherein R is from C₁-C₄ alkyl, which comprises cultivating a BU-4164E A or B-producing strain of *Streptomyces* sp. or mutant or variant thereof, and then recovering the BU-4164E A or B from the culture medium.

2. The process of Claim 1 wherein R is methyl or ethyl.
3. A biologically pure culture of the microorganism Streptomyces sp., ATCC 55001 or a variant or mutant thereof.
- 5 4. A process for preparing a pharmaceutical composition which comprises admixing an effective amount of at least one compound as defined in claims 1 or 2 with an inert pharmaceutically acceptable carrier or diluent.
- 10 5. Streptomyces sp., ATCC 55001 or variant or mutant thereof.

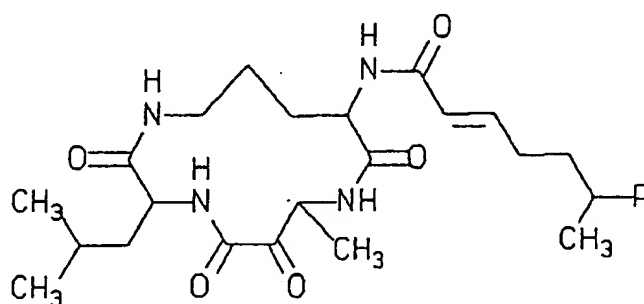
Claims for the following Contracting State: GR

1. A compound having the structure

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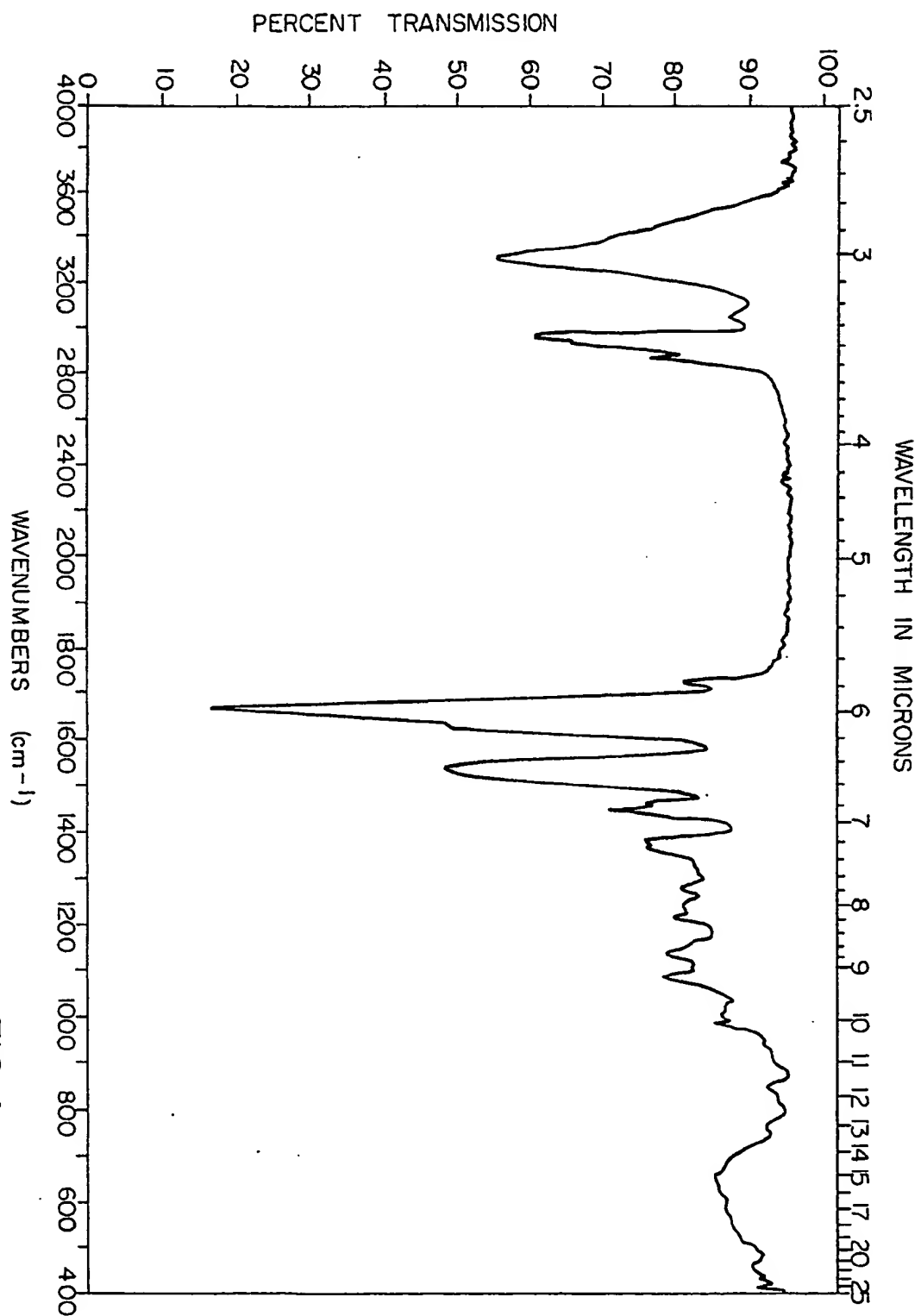
wherein R is from C₁-C₄ alkyl.

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2. The compound of Claim 1 wherein R is methyl or ethyl.
3. A process for producing a compound of claims 1 or 2 which comprises cultivating a BU-4164E A or B-producing strain of Streptomyces sp. or mutant or variant thereof, under submerged aerobic conditions in an aqueous nutrient medium containing assimilable sources of carbon and nitrogen until a substantial amount of BU-4164E A or B is produced by said organism in said culture medium and then recovering the BU-4164E A or B from the culture medium.
- 35 4. A biologically pure culture of the microorganism Streptomyces sp., ATCC 55001 or a variant or mutant thereof.
- 40 5. A process for preparing a pharmaceutical composition which comprises admixing an effective amount of at least one compound as defined in claims 1 or 2 with an inert pharmaceutically acceptable carrier or diluent.
- 45 6. Streptomyces sp., ATCC 55001 or variant or mutant thereof.

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FIG. 1

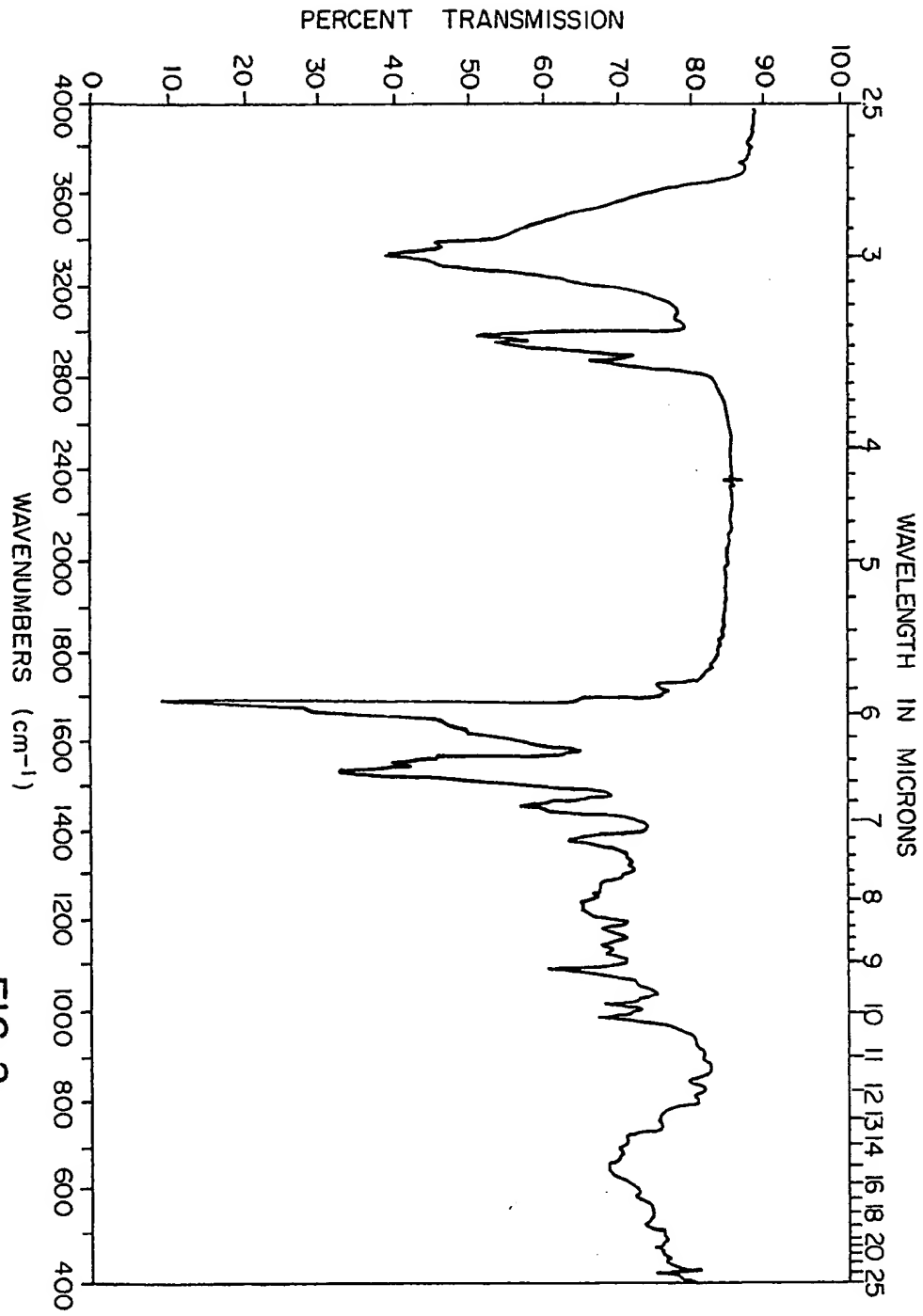


FIG. 2

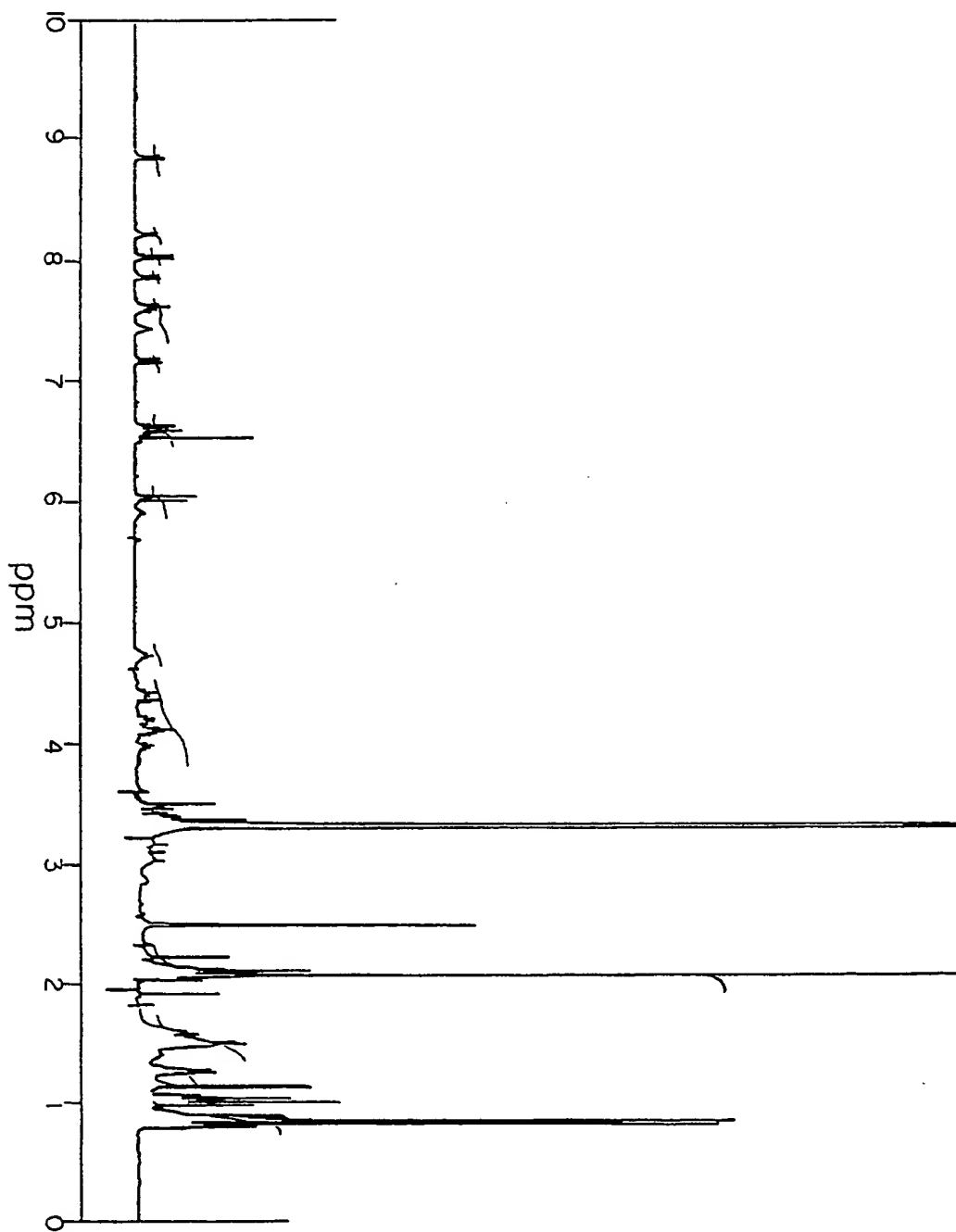


FIG. 3

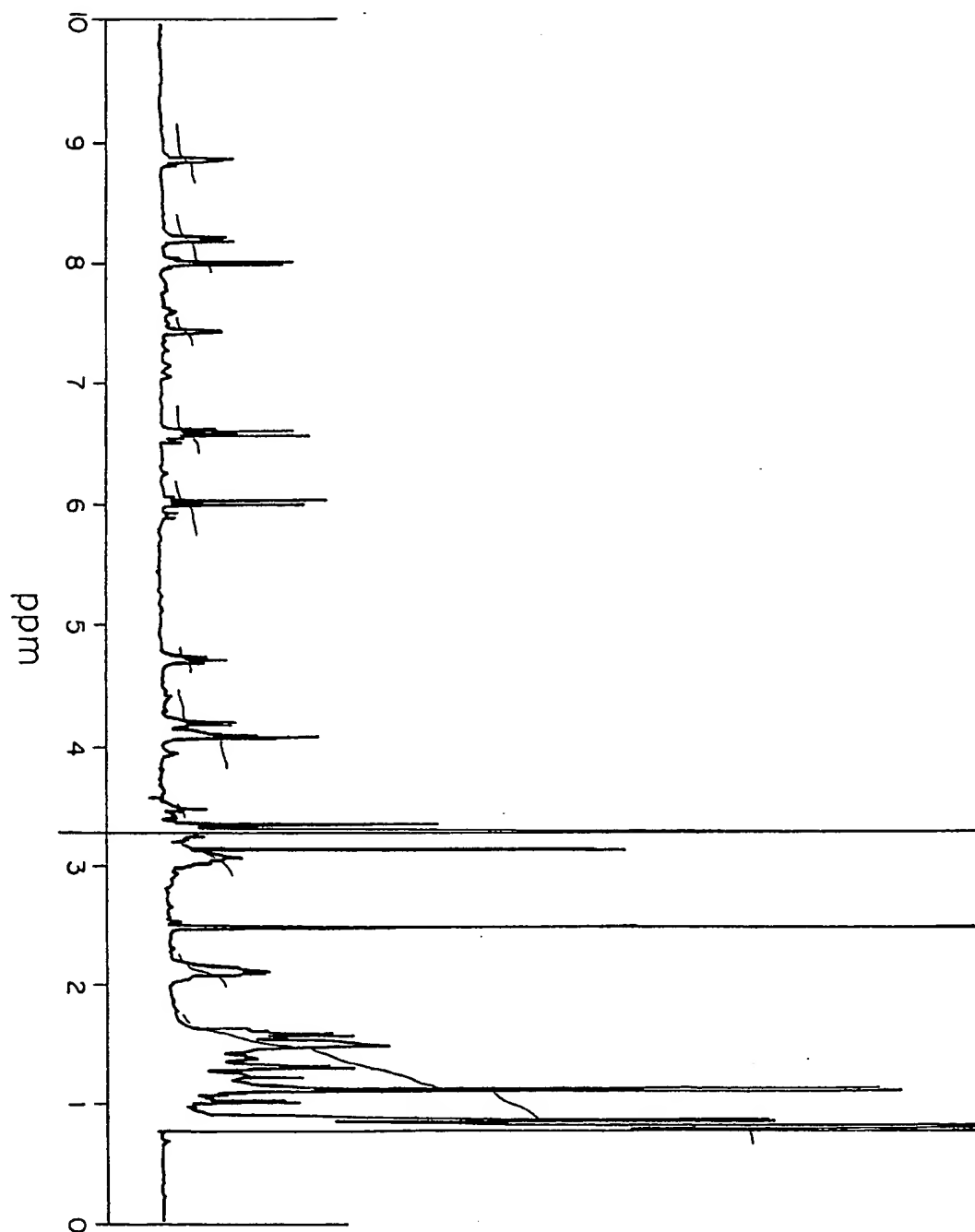
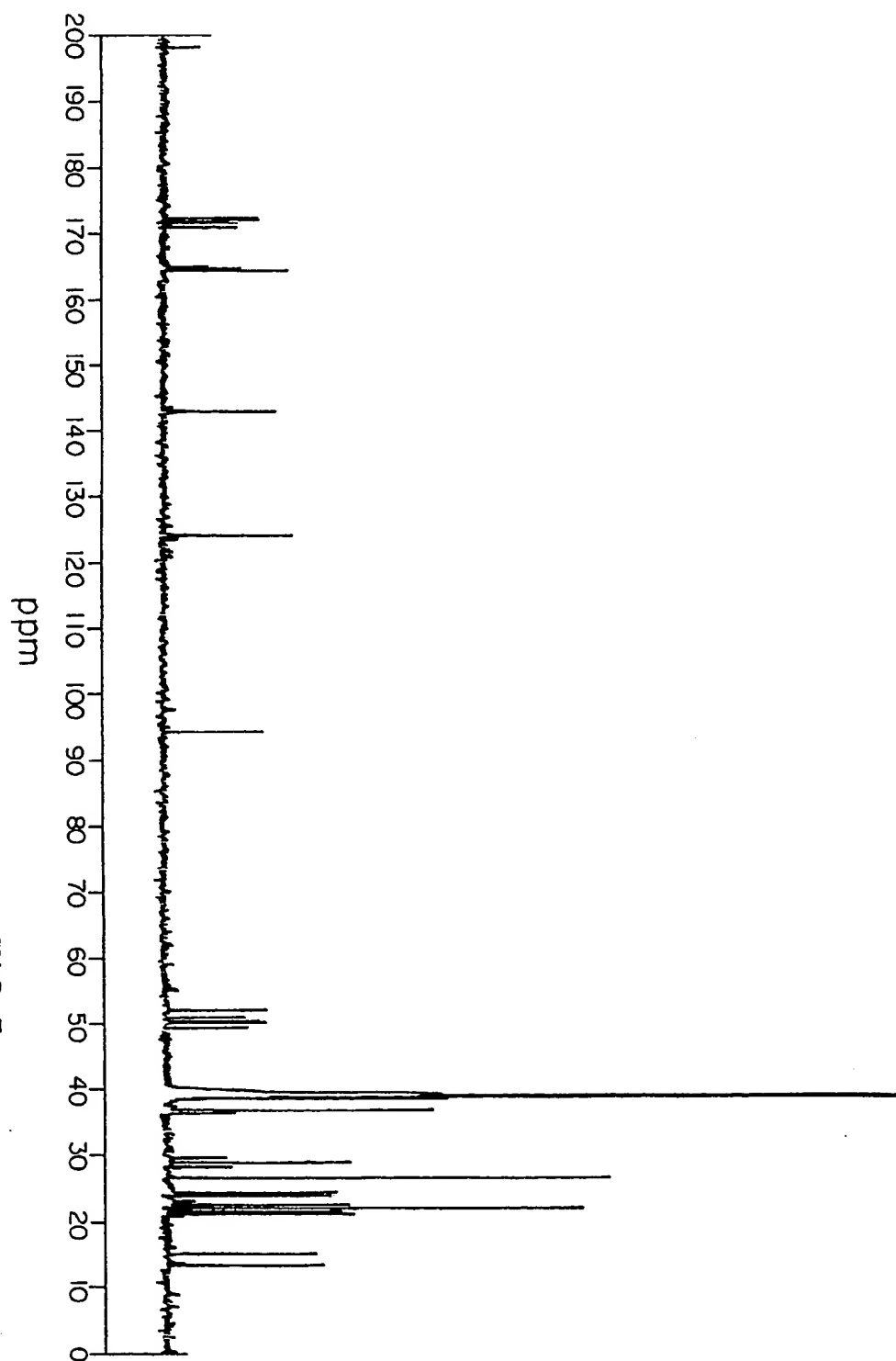


FIG. 4

FIG. 5

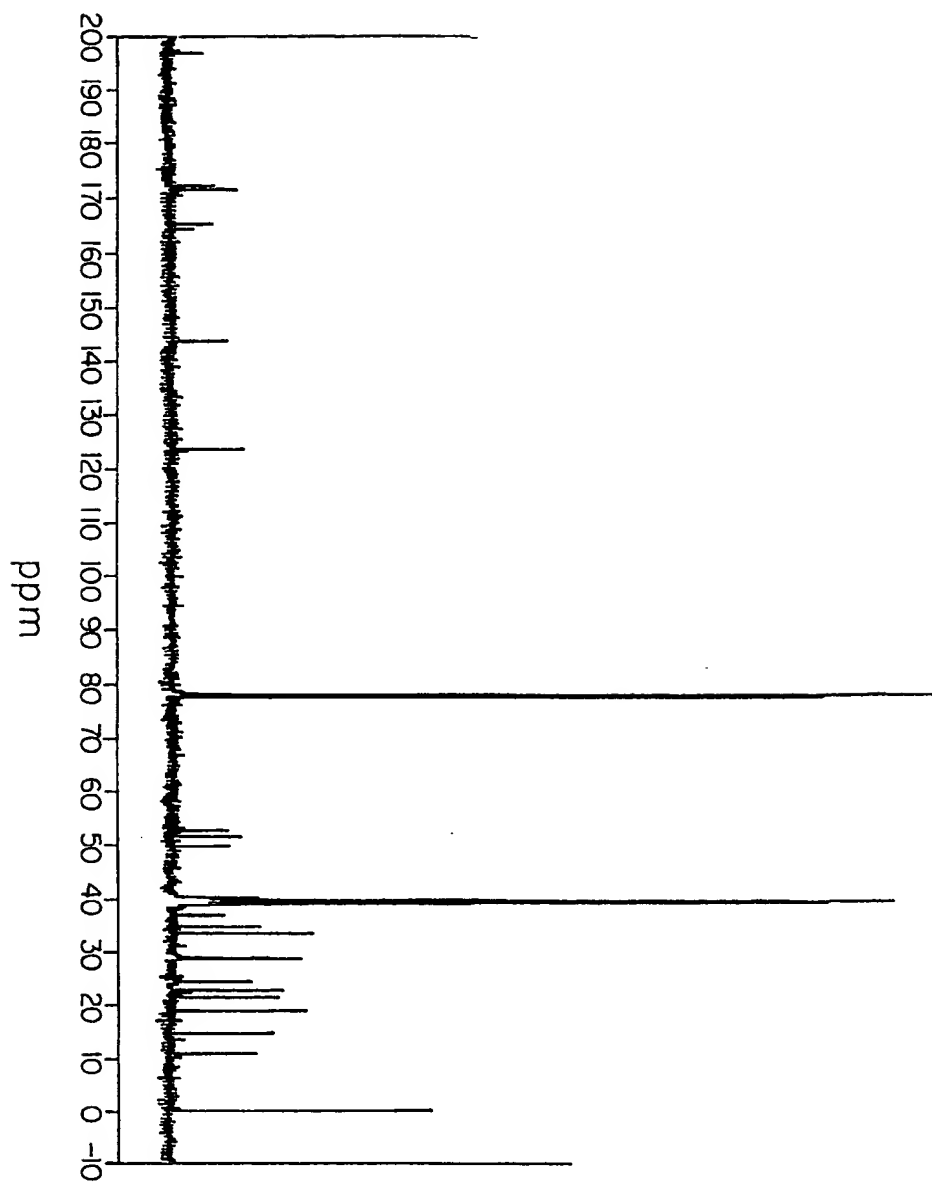


FIG. 6



European
Patent Office

EUROPEAN SEARCH REPORT

Application Number

EP 91 10 4163

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A	CHEMICAL ABSTRACTS, vol. 106, no. 25, 22nd June 1987, page 710, abstract no. 214388p, Columbus, Ohio, US; & JP-A-61 183 297 (SUNTORY LTD) 15-08-1986 - - - - -		C 07 D 255/02 A 61 K 31/395 C 12 P 17/10 C 12 N 1/20 // (C 12 P 17/10 C 12 R 1:465) (C 12 N 1/20 C 12 R 1:465)
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C 07 D 255/00 C 12 P 17/00
The present search report has been drawn up for all claims			
Place of search		Date of completion of search	Examiner
The Hague		24 June 91	ALLARD M.S.
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